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## **Familial Alzheimer's disease mutations at position 22 of the amyloid -peptide sequence differentially affect synaptic loss, tau phosphorylation and neuronal cell death in an ex vivo system**

Tackenberg, Christian ; Kulic, Luka ; Nitsch, Roger M

**Abstract:** Familial forms of Alzheimer's disease (AD) are caused by mutations in the presenilin genes or in the gene encoding for the amyloid precursor protein (APP). Proteolytic cleavage of APP generates the -amyloid peptide (A<sub>β</sub>), which aggregates into amyloid plaques, one of the major hallmarks of AD. APP mutations within the A<sub>β</sub> sequence, so-called intra-A<sub>β</sub> mutations, cluster around position E693 of APP, which corresponds to position E22 in the A<sub>β</sub> sequence. One of these mutations is the Osaka mutation, E693Δ, which has unique aggregation properties with patients showing unusually low brain amyloid levels on amyloid PET scans. Despite intense research on the pathomechanisms of different intra-A<sub>β</sub> mutants, our knowledge is limited due to controversial findings in various studies. Here, we investigated in an ex vivo experimental system the neuro- and synaptotoxic properties of two intra-A<sub>β</sub> mutants with different intrinsic aggregation propensities, the Osaka mutation E22Δ and the Arctic mutation E22G, and compared them to wild-type (wt) A<sub>β</sub>. Experiments in hippocampal slice cultures from transgenic mice were complemented by treating wild-type slices with recombinantly produced A<sub>β</sub> 40 or A<sub>β</sub> 42 containing the respective intra-A<sub>β</sub> mutations. Our analyses revealed that wt A<sub>β</sub> and E22G A<sub>β</sub>, both recombinant and transgenic, caused a loss of dendritic spines along with an increase in tau phosphorylation and tau-dependent neurodegeneration. In all experiments, the 42-residue variants of wt and E22G A<sub>β</sub> showed stronger effects than the respective A<sub>β</sub> 40 isoforms. In contrast, E22Δ A<sub>β</sub> neither reduced dendritic spine density nor resulted in increased tau phosphorylation or neuronal cell death in our ex vivo system. Our findings suggest that the previously reported major differences in the aggregation kinetics between E22G and E22Δ A<sub>β</sub> are likely reflected in different disease pathomechanisms.

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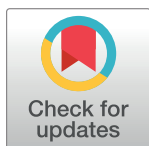
RESEARCH ARTICLE

# Familial Alzheimer's disease mutations at position 22 of the amyloid $\beta$ -peptide sequence differentially affect synaptic loss, tau phosphorylation and neuronal cell death in an *ex vivo* system

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## Abstract

Familial forms of Alzheimer's disease (AD) are caused by mutations in the presenilin genes or in the gene encoding for the amyloid precursor protein (APP). Proteolytic cleavage of APP generates the  $\beta$ -amyloid peptide ( $A\beta$ ), which aggregates into amyloid plaques, one of the major hallmarks of AD. APP mutations within the  $A\beta$  sequence, so-called intra- $A\beta$  mutations, cluster around position E693 of APP, which corresponds to position E22 in the  $A\beta$  sequence. One of these mutations is the Osaka mutation, E693 $\Delta$ , which has unique aggregation properties with patients showing unusually low brain amyloid levels on amyloid PET scans. Despite intense research on the pathomechanisms of different intra- $A\beta$  mutants, our knowledge is limited due to controversial findings in various studies. Here, we investigated in an *ex vivo* experimental system the neuro- and synaptotoxic properties of two intra- $A\beta$  mutants with different intrinsic aggregation propensities, the Osaka mutation E22 $\Delta$  and the Arctic mutation E22G, and compared them to wild-type (wt)  $A\beta$ . Experiments in hippocampal slice cultures from transgenic mice were complemented by treating wild-type slices with recombinantly produced  $A\beta$ 40 or  $A\beta$ 42 containing the respective intra- $A\beta$  mutations. Our analyses revealed that wt  $A\beta$  and E22G  $A\beta$ , both recombinant and transgenic, caused a loss of dendritic spines along with an increase in tau phosphorylation and tau-dependent neurodegeneration. In all experiments, the 42-residue variants of wt and E22G  $A\beta$  showed stronger effects than the respective  $A\beta$ 40 isoforms. In contrast, E22 $\Delta$   $A\beta$  neither reduced dendritic spine density nor resulted in increased tau phosphorylation or neuronal cell death in our *ex vivo* system. Our findings suggest that the previously reported major differences in the aggregation kinetics between E22G and E22 $\Delta$   $A\beta$  are likely reflected in different disease pathomechanisms.

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## Introduction

Alzheimer's Disease (AD) is the most common age-related neurodegenerative disorder. The major histopathological hallmarks are the presence of extracellular aggregates of A $\beta$  in amyloid plaques and intracellular neurofibrillary tangles containing hyperphosphorylated tau as major component. According to the amyloid hypothesis A $\beta$  aggregates, especially oligomers, are the driver of AD pathology [1]. A $\beta$  is generated by proteolytic processing of APP by the  $\beta$ - and  $\gamma$ -secretase complexes. Familial forms of AD are caused by mutations in presenilin 1 or presenilin 2, the catalytic subunits of the  $\gamma$ -secretase complex, or by mutations in APP itself. APP mutations can have various effect on APP processing, depending on their location. Mutations around the  $\beta$ -secretase cleavage site, such as the APP<sub>swe</sub> mutation K670N/M671L, increase the general production of A $\beta$ , whereas mutations near the  $\gamma$ -secretase cleavage site shift the processing towards longer isoforms of A $\beta$ , such as A $\beta$ 42, which is thought to be more toxic than the shorter A $\beta$ 40 variant [2]. Mutations within the A $\beta$  sequence of APP alter the aggregation propensity of A $\beta$ . A special cluster of these intra-A $\beta$  mutations is localized at position E693 of APP, corresponding to position E22 in the A $\beta$  sequence, a hydrophobic core of the peptide. Mutations at this position include the Dutch (E693Q) [3], Italian (E693K) [4], Arctic (E693G) [5] or Osaka (E693 $\Delta$ ) mutation [6].

The Arctic mutation E693G was discovered in 2001 in a Swedish family showing clinical symptoms of early-onset AD. Carriers of this mutation show decreased A $\beta$ 42 and A $\beta$ 40 levels in plasma [5] and low PiB PET retention while cerebral glucose metabolism and CSF levels of A $\beta$ 42, total and phosphorylated tau are highly pathologic [7]. The Osaka mutation E693 $\Delta$ , discovered in 2008, shows a recessive pattern of inheritance that causes AD-like symptoms despite very low PiB PET signals [6]. As synthetic E22 $\Delta$  A $\beta$  was shown not to form fibrils, it was hypothesized that this mutation favours oligomerization causing AD in the absence of fibril/plaque formation [6]. However, the literature on the aggregation propensity of the E693 $\Delta$  mutation is controversial as subsequent studies, including our own research using recombinant A $\beta$  peptides, showed fibril formation by E22 $\Delta$  A $\beta$  [8–11]. Compared to wt and E22G, recombinant E22 $\Delta$  A $\beta$  40 and A $\beta$  42 had unique aggregation characteristics in thioflavin T aggregation assays with E22 $\Delta$  A $\beta$ 42 forming SDS-resistant amyloid fibrils direct after reconstitution, while fibril growth occurred non-exponentially. In contrast, E22 $\Delta$  A $\beta$ 40 aggregation was characterized by an exponential growth phase [8]. These findings were supported *in vivo*. Extracellular accumulation of insoluble A $\beta$  occurred in E22 $\Delta$ A $\beta$  transgenic mice as leptomeningeal cerebral amyloid angiopathy at the age of 24 months, while intraneuronal A $\beta$  oligomers were detected already at 3 months. In contrast, arcA $\beta$  transgenic mice, expressing E22G A $\beta$  and Tg2576 mice, which express wt A $\beta$ , accumulated detergent-insoluble, fibrillar aggregates already at 6 month (arcA $\beta$  mice) and 15 months (Tg2576) of age [8].

Besides several studies on the mechanisms of E22 $\Delta$  A $\beta$  toxicity, a direct comparison to wt and E22G A $\beta$  for its neuro- and synaptotoxic properties has not been performed yet. Therefore, we analyzed the effects of recombinant E22 $\Delta$ , E22G and wt A $\beta$ 40 and A $\beta$ 42 on neuronal survival and dendritic spine densities in an *ex vivo* experimental paradigm in hippocampal slice cultures. Treatment of wild-type slices with recombinant A $\beta$  was directly compared to slices obtained from the respective transgenic mouse models.

We found that both recombinant wt and E22G A $\beta$ 42 induced loss of dendritic spines, increased tau phosphorylation and caused a tau-dependent cell death in wild-type slices. This was confirmed in slices from Tg2576 and arcA $\beta$  transgenic mice. In contrast, neither induction of toxicity nor an increase in tau phosphorylation were observed in slices treated with E22 $\Delta$  A $\beta$ 40 or A $\beta$ 42 or in slices from E22 $\Delta$ A $\beta$  transgenic mice, suggesting differential disease pathomechanisms for distinct intra-A $\beta$  mutants.

Table 1. APP/A $\beta$  mutants used in this study.

	Tg2576 mice [16]	arcA $\beta$ mice [12]	E22 $\Delta$ A $\beta$ mice [8]	recombinant wt A $\beta$	recombinant E22G A $\beta$	recombinant E22 $\Delta$ A $\beta$
APP/A $\beta$ mutations	APP <sub>swe</sub> (K670N + M671L)	APP <sub>swe</sub> (K670N + M671L); APP <sub>arc</sub> (E693G (= E22G))	APP <sub>swe</sub> (K670N + M671L); APP <sub>osaka</sub> (E693 $\Delta$ (= E22 $\Delta$ ))	no mutation	Arctic mutation; E22G (= E693G)	Osaka mutation; E22 $\Delta$ (= E693 $\Delta$ )

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## Materials and methods

### Animals

ArcA $\beta$  mice and E22 $\Delta$ A $\beta$  mice were obtained as previously described [8, 12]. Tg2576 mice (B6;SJL-Tg(APP<sub>SWE</sub>)2576Kha) were obtained from Taconic Biosciences, Rensselaer, NY, USA. Details on respective APP mutations are displayed in Table 1. All animal experiments were performed in compliance with Swiss national guidelines and were approved by the veterinarian office of the Canton of Zurich. Animals were housed in groups. For preparation of hippocampal slice cultures P6-P8 mice were sacrificed by decapitation.

### Organotypic hippocampal slice cultures

Organotypic hippocampal slice cultures were prepared from P6-P8 mice and cultured as previously described [13], with 3 slices per well. For assessment of dendritic spine density, cultures were transduced with Sindbis virus expressing EGFP at DIV 12 and were fixed at DIV15 with 4% paraformaldehyde in PBS containing 4% sucrose for 2h at 4°C. After washing with PBS, cultures were mounted with Hydromount (National diagnostics, Atlanta, Georgia) and cover-slipped. For analysis of hTau-dependent toxicity, slices were transduced at DIV 12 with Sindbis virus expressing EGFP-coupled human 441 wild-type tau. At DIV 16 culture medium was harvested for cytotoxicity assays and lysates were prepared for western blot analyses. To determine the effect of recombinant A $\beta$  on dendritic spines, slices were treated with 1  $\mu$ M recombinant A $\beta$  from DIV 8 until fixation on DIV 15. For measurement of cell toxicity, 1  $\mu$ M recombinant A $\beta$  was applied from DIV 12 until DIV 16. This concentration was chosen as it was previously shown that 1  $\mu$ M of recombinant wt A $\beta$  induces significant loss of dendritic spines and a human tau-dependent neuronal cell death in this experimental system [13].

### Recombinant A $\beta$

Recombinant A $\beta$  was prepared as described previously [8, 9, 14]. For reconstitution, lyophilized A $\beta$  was dissolved in 10 mM NaOH and sonicated for 1 min. Concentration was confirmed using Nanodrop (Thermo Scientific) and if necessary, adjusted to the concentration of 62.5  $\mu$ M using 10 mM NaOH. Before addition to cell culture medium, the A $\beta$  solution was mixed in neutralization buffer (40 mM HCl, 50mM H<sub>3</sub>PO<sub>4</sub>-NaOH pH 7.4, 10 mM NaCl) in a 4:1 ratio yielding a 50  $\mu$ M A $\beta$  solution at pH7.4 which is immediately added to the cell culture in a 1:50 ratio. For all steps, LoBind tubes (Eppendorf, Germany) were used.

### Analysis of dendritic spine density

Apical dendritic segments length in CA1 stratum radiatum were imaged using Leica SP2 confocal laserscanning microscope, equipped with 63x objective (NA: 1.2) and 488nm Argon laser. CA1 apical dendrites were chosen for their straighter appearance compared to CA3 dendrites, which allows the imaging of a single dendritic segment and a less error-prone counting of spine numbers per dendrite. Further, it has been shown that A $\beta$  affects spine numbers in CA1 and CA3 apical dendrites to a similar extent [15]. Images were taken at a size of 30x30  $\mu$ m

(512x512 pixel, voxel size: 0.05813x0.05813x0.25  $\mu$ m) with one clearly visible dendritic segment of 30–40  $\mu$ m length. Image stacks were processed to maximum projections, regions of interest were cropped if applicable and dendritic spine density was determined as spine counts per  $\mu$ m dendrite using ImageJ.

### Assessment of cell death

To determine cytotoxicity of A $\beta$ , culture medium was harvested on DIV 16, frozen in liquid nitrogen and stored at -80°C until further analysis with CytotoxGlo assay (Promega, Madison, USA), according to the manufacturer's recommendations. For measurement of human EGFP-tau-dependent toxicity, CytotoxGlo signals were normalized to EGFP fluorescence using microplate reader Synergy HT.

### Westernblot analysis

On DIV 16 slices were harvested in RIPA buffer (50 mM Tris-HCl, 150 mM NaCl, 2 mM EDTA, 1% NP-40, 0.5% deoxycholate, and 0.1% SDS, pH 8.0) containing phosphatase inhibitor cocktails 1 and 2 (Sigma) and protease inhibitor cocktail (Roche) and centrifuged at 5000xg for 10 min at 4°C. The supernatant was collected, frozen in liquid nitrogen and stored at -80°C. Samples were mixed with loading buffer, resolved using 10–20% SDS PAGE and transferred to nitrocellulose membranes (Millipore). Immunoblotting was performed using primary antibodies Anti-GFP to detect total EGFP-coupled tau (1:1000, Roche), anti phospho-Tau AT8 (1:200, Pierce), anti GAPDH (1:5000, Biodesign) and HRP-conjugated secondary antibody (1:2000, Amersham). Bands were detected using the Supersignal Femto Maximum Sensitivity Substrate (Thermo) and imaged with Fujifilm Las-3000 assuring that no pixels were saturated. Band intensities were quantified with ImageJ and corrected for background.

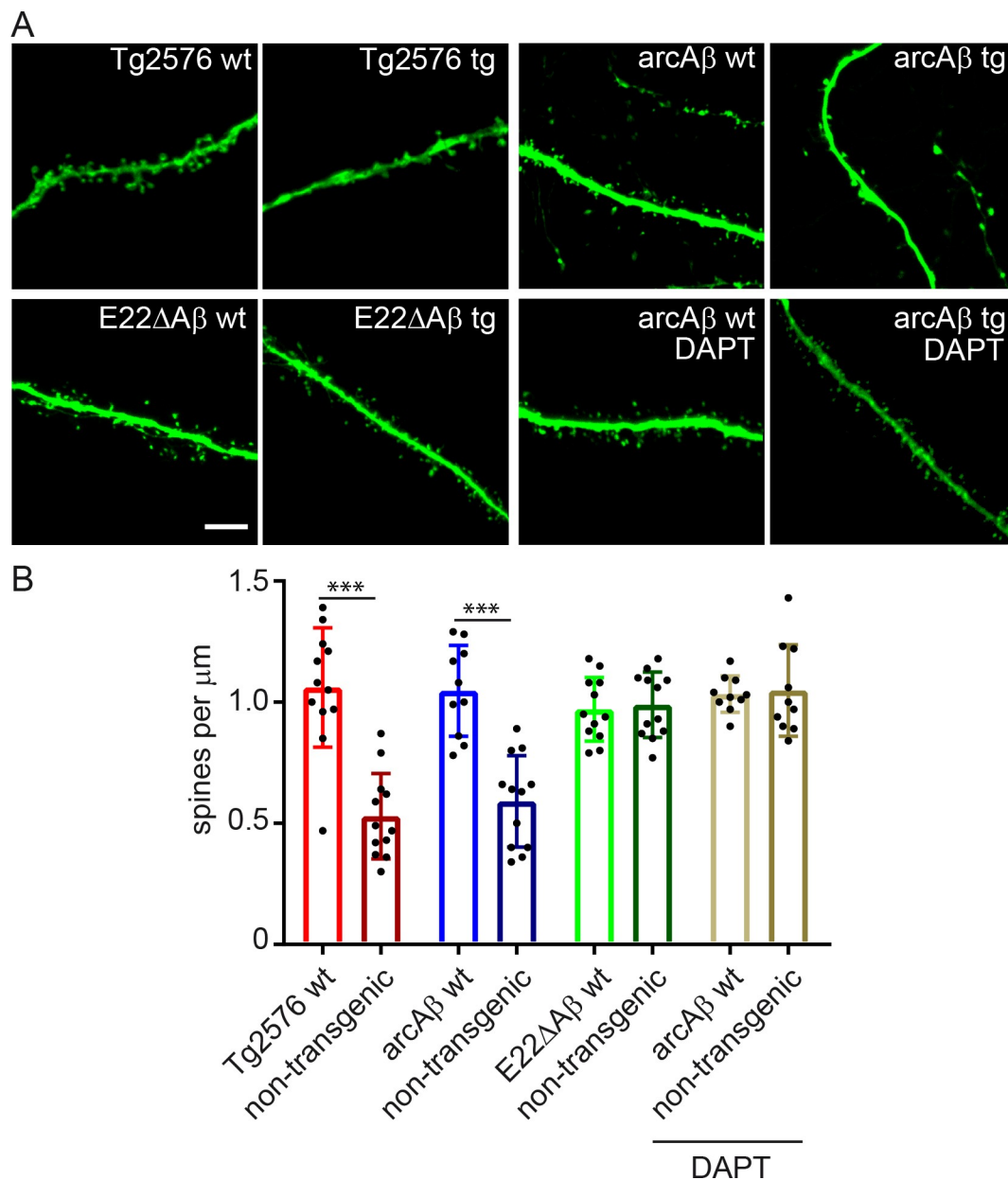
### Statistics

Data are presented as mean  $\pm$  standard deviation (SD). All datapoints (n-numbers) are plotted individually in each bar graph. Statistical analysis was performed with GraphPad Prism 7.01 software using Shapiro-Wilk test for normal distribution followed by one-way ANOVA with Tukey's multiple comparison test. Exact p-values, mean differences and significances for all experiments are listed in [S1 Table](#). For dendritic spine analysis, n = 1 refers to one image containing one dendritic segment. For cell death assessment, n = 1 represents the analysis of supernatant of one cell culture well in which 3 hippocampal slices were cultured. For western blot analysis, n = 1 refers to the lysate of one well, in which 3 hippocampal slices were cultured and pooled for analysis.

## Results

### Reduced dendritic spine density in slice cultures from Tg2576 and arcA $\beta$ -but not E22 $\Delta$ A $\beta$ transgenic mice

Loss of pre- and postsynaptic structures is an early feature of AD and directly correlates to the degree of dementia [17]. Whether different intra-A $\beta$  mutations confer a different degree of synaptotoxicity to A $\beta$  is largely unknown. To determine the differential mechanisms of synapse loss caused by intra-A $\beta$  mutants, we applied the well-established organotypic hippocampal slice culture technique [13, 15, 18, 19]. Slices were obtained from i) Tg2576 transgenic mice, which do not have an intra-A $\beta$  mutation thus producing wild-type A $\beta$ , ii) arcA $\beta$  transgenic mice, producing A $\beta$  with the "Arctic" mutation E22G and iii) E22 $\Delta$ A $\beta$  transgenic mice with the "Osaka" mutation E22 $\Delta$  ([Table 1](#)). Slices were transduced with EGFP-expressing



**Fig 1. Reduction in dendritic spine density in hippocampal slices from Tg2576 and arcA $\beta$ - but not E22 $\Delta$ A $\beta$  transgenic mice.** A: Representative confocal images of dendritic segments of hippocampal slices from Tg2576, arcA $\beta$ - and E22 $\Delta$ A $\beta$  transgenic mice and the respective non-transgenic littermates. B: Spine counts per  $\mu$ m dendrite. Spine density was reduced in cultures from Tg2576 tg mice and arcA $\beta$  tg mice but not in E22 $\Delta$ A $\beta$  tg mice. Treatment with 0.5  $\mu$ M DAPT abolished spine reduction in arcA $\beta$  tg slices.  $n = 10$ – $12$ ; Data are means  $\pm$  SD. Statistical significance was assessed by one-way ANOVA with Tukey's multiple comparison test (\*\* $p < 0.001$ ). Scale bar: 5  $\mu$ m.

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Sindbis virus, allowing high-resolution imaging of dendritic segments and spines [19]. As shown previously [13], spine density was strongly reduced in slices from arcA $\beta$  mice compared to slices from non-transgenic (non-tg) littermates (Fig 1A and 1B). This was rescued by treatment with the  $\gamma$ -secretase inhibitor DAPT at 0.5  $\mu$ M indicating that spine reduction is mediated by A $\beta$  and showing the validity of the transgenic slice culture model. Slices from Tg2576 mice displayed a similar reduction in spine numbers compared to arcA $\beta$  tg mice whereas



E22 $\Delta$ A $\beta$  tg slices showed no differences to their non-tg control (E22 $\Delta$ A $\beta$  wt). A detailed list of all p-values is presented in [S1 Table](#).

### Recombinant wt and E22G A $\beta$ but not E22 $\Delta$ A $\beta$ reduce dendritic spine density

To confirm the results obtained in slices from the transgenic mice, we used wild-type slices treated with recombinant A $\beta$  containing the respective intra-A $\beta$  mutations. Recombinant A $\beta$  was produced in *Escherichia coli* with higher purity than commercially available synthetic A $\beta$  [14]. Wt, E22G and E22 $\Delta$  A $\beta$  were tested at a concentration of 1  $\mu$ M as 40- and 42-residue isoforms. Treatment with wt A $\beta$ 42 and E22G A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 42 induced a strong reduction in dendritic spine density compared to untreated and vehicle treated slices ([Fig 2A and 2B](#)). Interestingly, the degree of spine loss caused by wt A $\beta$ 42 and E22G A $\beta$ 42 was similar to the spine loss observed in slices from the respective transgenic mouse model ([Fig 1](#)). Wt A $\beta$ 40 and E22G A $\beta$ 40 caused spine loss to a lesser extent, with wt A $\beta$ 40 not reaching significance ( $p = 0.06$ ). E22 $\Delta$  A $\beta$ 40 did not show an effect on spine density. A direct comparison between the 40- and 42-residue isoforms showed that E22G A $\beta$ 42 reduced spine numbers significantly more than E22G A $\beta$ 40 ([S1A Fig](#)). While spine reduction upon wt A $\beta$ 42 treatment was similarly stronger than upon treatment with wt A $\beta$ 40 treatment, this difference did not reach significance ( $p = 0.059$ ). However, no difference was observed between E22 $\Delta$  A $\beta$ 42 and A $\beta$ 40 ( $p = 0.909$ ) ([S1A Fig](#)) with both isoforms showing no significant effect on dendritic spines.

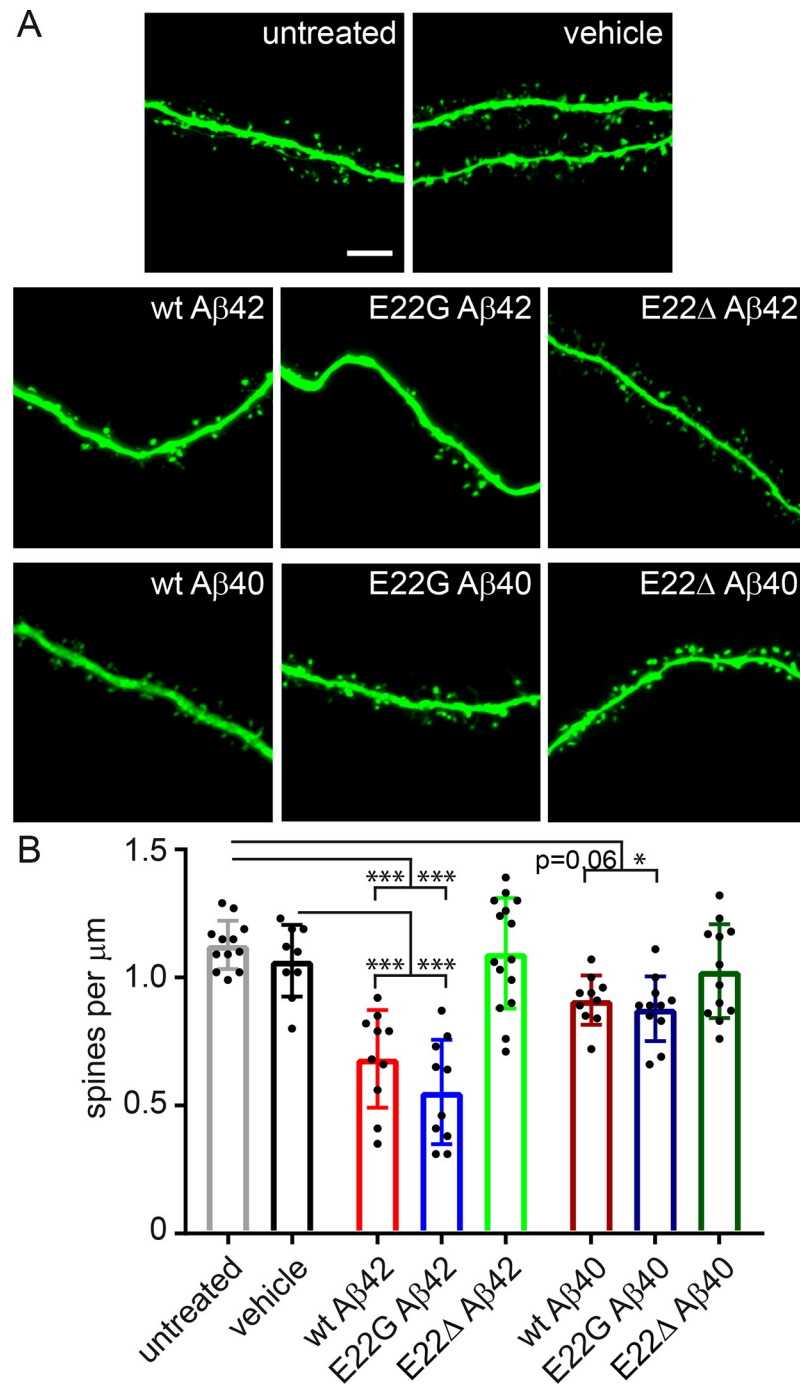
### Wt and E22G A $\beta$ but not E22 $\Delta$ A $\beta$ induce of tau phosphorylation and tau-dependent cell death

We and others have shown that A $\beta$  induces cell death, depending on the presence of tau [20], especially of human tau [15]. To determine whether intra-A $\beta$  mutations confer different levels of toxicity to A $\beta$ , we first used slices from Tg2576, arcA $\beta$  tg and E22 $\Delta$ A $\beta$  tg mice, and virally expressed the 441 isoform of human tau, coupled to EGFP. Toxicity was determined using CytotoxGlo assay, measuring dead-cell protease activity released from cells that have lost membrane integrity ([Fig 3A](#)). A significantly higher toxicity was observed in slices from Tg2576 mice compared non-tg littermates. Higher signals were also found in arcA $\beta$  tg compared to non-tg slices, although this did not reach significance ( $p = 0.058$ ). No difference was observed between slices from E22 $\Delta$ A $\beta$  tg and non-tg littermates ( $p = 0.843$ ).

To support this finding, slices from wild-type mice expressing human tau were treated with 1  $\mu$ M recombinant A $\beta$  ([Fig 3B](#)). Similar to the findings in transgenic cultures, wt A $\beta$ 42 and E22G A $\beta$ 42 treatment increased toxicity compared to untreated and vehicle treated slices. Wt A $\beta$ 40 and E22G A $\beta$ 40 did not induce toxicity. E22 $\Delta$  A $\beta$ 42 and E22 $\Delta$  A $\beta$ 40 both did not affect cell viability in this setting, confirming the results from E22 $\Delta$ A $\beta$  tg cultures. A direct comparison between the 40- and 42-residue isoforms showed that wt A $\beta$ 42 and E22G A $\beta$ 42 were significantly more toxic than wt A $\beta$ 40 and E22G A $\beta$ 40, respectively, whereas no difference was observed between E22 $\Delta$  A $\beta$ 42 and E22 $\Delta$  A $\beta$ 40 ( $p = 0.688$ ) ([S1B Fig](#)). To further confirm that A $\beta$  toxicity is mediated by the presence of human tau, wild-type slices were treated with 1  $\mu$ M recombinant A $\beta$  in the absence of human tau expression ([Fig 3C](#)). In this case, no A $\beta$  toxicity was observed.

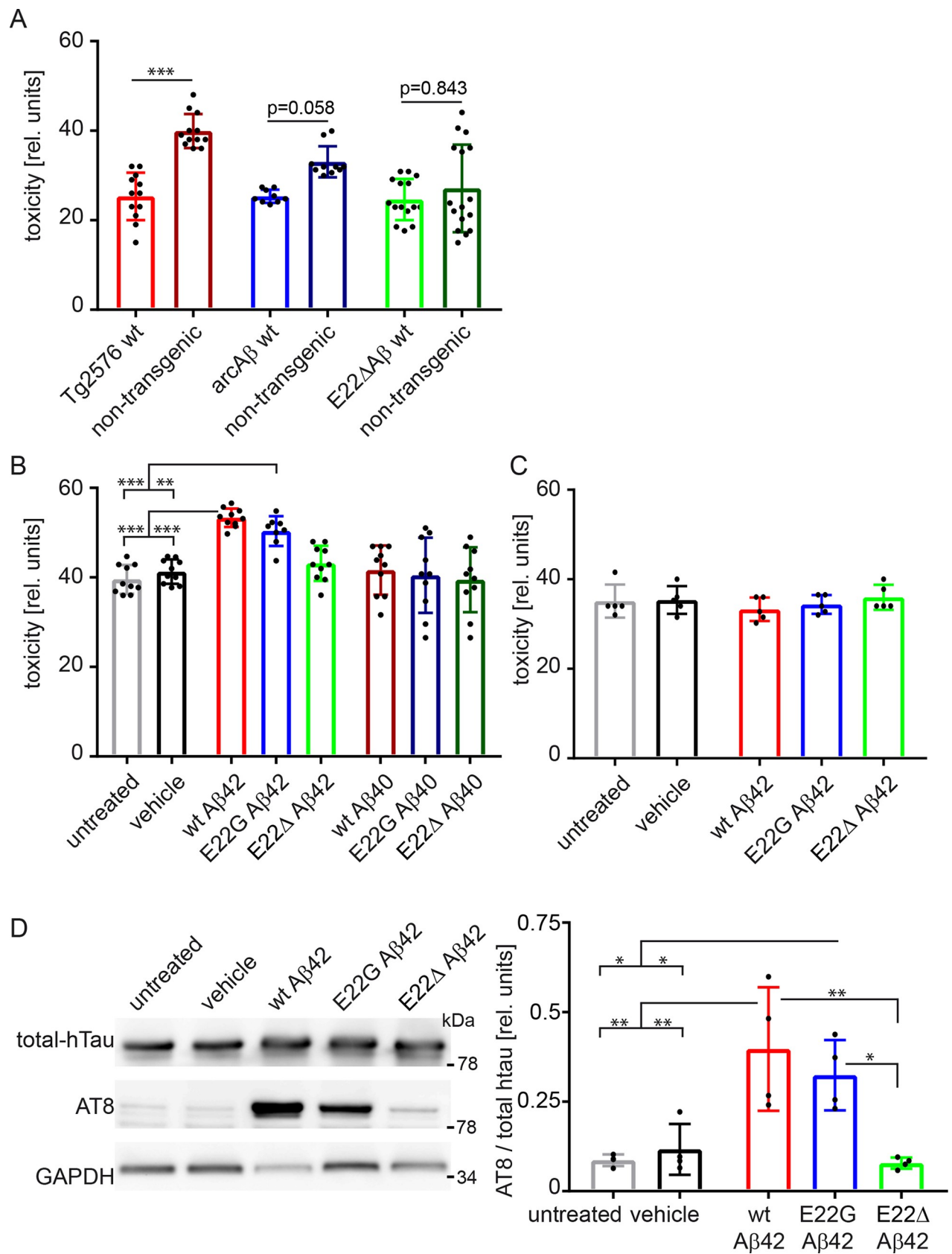
It has been shown that A $\beta$  can activate tau kinases thereby increasing tau phosphorylation which induces cell death downstream of A $\beta$  [15, 21]. We determined the level of tau phosphorylation at the AT8 epitope, in cultures treated with wt A $\beta$ 42, E22G A $\beta$ 42 or E22 $\Delta$  A $\beta$ 42 ([Fig 3D](#)). In line with our findings on cell toxicity, only wt A $\beta$ 42 and E22G A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 42 significantly increased tau phosphorylation.





**Fig 2. Reduction in dendritic spine density in hippocampal slices treated with recombinant wt and E22G A $\beta$  but not E22 $\Delta$  A $\beta$ .** A: Representative confocal images of dendritic segments of wild-type mice treated with 1 μM recombinant A $\beta$  for 7 days. B: Spine counts per μm dendrite. Spine density was reduced after treatment with wt A $\beta$ 42 and E22G A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 42. Incubation with wt A $\beta$ 40 and E22G A $\beta$ 40 caused a mild reduction in spine numbers, whereas no effect was observed for E22 $\Delta$  A $\beta$ 40. n = 10–15; Data are means  $\pm$  SD. Statistical significance was assessed by one-way ANOVA with Tukey's multiple comparison test (\*p < 0.05; \*\*\*p < 0.001). Scale bar: 5 μm.

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**Fig 3. Induction of tau phosphorylation and tau-dependent cell death by wt and E22G A $\beta$  but not E22 $\Delta$  A $\beta$ .** A: Cytotoxicity in human tau expressing slices from Tg2576, arcA $\beta$ - and E22 $\Delta$ A $\beta$  tg mice and the respective non-transgenic littermates, measured with CytotoxGlo assay. Increased cell death was observed in Tg2576 and arcA $\beta$  tg slices, although not significant in arcA $\beta$  tg slices, but not in E22 $\Delta$ A $\beta$  tg slices.  $n = 9-16$ . B: Cytotoxicity in human tau expressing slices from wild-type mice treated with 1  $\mu$ M recombinant A $\beta$ . Treatment with wt A $\beta$ 42 and E22G A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 42, wt A $\beta$ 40 or E22 $\Delta$  A $\beta$ 40 increased cell death.  $n = 8-10$ . C: Cytotoxicity in slices from wild-type mice treated with 1  $\mu$ M recombinant A $\beta$ , in the absence of human tau expression. No toxicity was observed by wt A $\beta$ 42, E22G A $\beta$ 42 or E22 $\Delta$  A $\beta$ 42 in the absence of human tau.  $n = 5$ . D: Representative western blot (left) of lysate of human tau expressing wild-type slices, treated with 1  $\mu$ M recombinant A $\beta$ , displaying total human tau, phospho-tau at the AT8 epitope and GAPDH. Quantification (right) of western blots showed increased tau phosphorylation after treatment with wt A $\beta$ 42 and E22G A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 42.  $n = 4$ . Data are means  $\pm$  SD. Statistical significance was assessed by one-way ANOVA with Tukey's multiple comparison test (\* $p < 0.05$ ; \*\* $p < 0.01$ , \*\*\* $p < 0.001$ ).

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Taken together, wt and E22G A $\beta$ —either recombinant or transgenic—but not E22 $\Delta$  A $\beta$  increased tau phosphorylation and tau-dependent cell death.

## Discussion

In this study, we showed that the Osaka intra-A $\beta$  mutation E693 $\Delta$  did not confer toxic properties to A $\beta$  in an ex vivo tissue culture model, compared to the Arctic intra-A $\beta$  mutation E693G and to wt A $\beta$ . We used either wild-type organotypic hippocampal slices treated with recombinant preparations of wt, E22G or E22 $\Delta$  A $\beta$ 42 or A $\beta$ 40 or slices cultured from the respective transgenic mouse model, i.e. Tg2576, arcA $\beta$  and E22 $\Delta$ A $\beta$  mice (Table 1). All mouse lines express APP<sub>swc</sub> under the control of the prion protein promoter and differ only in the intra-A $\beta$  mutation at position 22. Expression levels of APP are identical in these mice [8]. Several studies on E22 $\Delta$  aggregation or toxicity used synthetic A $\beta$  peptides. However, synthetically produced A $\beta$  is often contaminated with up to 9% impurities, such as racemized peptides, that cannot be avoided due to the technical limitations of peptide synthesis [9, 14]. Importantly, impurities in synthetic A $\beta$  already in the range of 3% were previously shown to have a significant impact on its aggregation properties [9, 14]. The use of recombinant A $\beta$  can bypass this limitation and it has been shown that recombinantly produced A $\beta$  aggregates faster and is more toxic than synthetic A $\beta$  [14]. In a previous study, we compared the aggregation propensity of recombinant wt, E22G and E22 $\Delta$  A $\beta$ 42 and A $\beta$ 40 and found unique aggregation characteristics of E22 $\Delta$  A $\beta$ 42 and A $\beta$ 40 [8]. Compared to E22G and wt A $\beta$ , E22 $\Delta$  A $\beta$ 42 displayed highest fibrillogenesis propensity as it aggregated without a lag phase in a thioflavin T assay and formed SDS-stable high molecular weight aggregates directly after reconstitution whereas fibril growth after reconstitution occurred slowly and non-exponentially. Thus, addition of recombinant E22 $\Delta$  A $\beta$  to our slice cultures may have caused rapid formation of less toxic fibrils in the extracellular space, which may explain the lack of toxicity in this setting (see also discussion below).

Loss of pre- and postsynaptic contacts is a major correlate to the degree of dementia in AD [17]. We showed that recombinant wt A $\beta$ 42 and E22G A $\beta$ 42 strongly reduced the numbers of dendritic spines in slice cultures while wt A $\beta$ 40 and E22G A $\beta$ 40 only had a mild effect. As A $\beta$  treatment in the absence of human tau expression is not toxic in our model, the reduction in dendritic spine density represents a distinct pathophysiological event caused by A $\beta$  rather than a general toxic effect. Importantly, the degree of spine loss in the A $\beta$ 42-treated wild-type slices was similar to spine loss in slices from the respective transgenic mice, confirming that the treatment with recombinant A $\beta$  can mirror the effects of A $\beta$  produced from processing of APP. Further, the protective effect of  $\gamma$ -secretase inhibitor DAPT indicates that the synaptotoxic events in these transgenic cultures were indeed caused by A $\beta$ . In contrast to wt and E22G A $\beta$ , dendritic spine loss was neither observed in wild-type slices treated with recombinant E22 $\Delta$  A $\beta$ 42 or A $\beta$ 40 nor in slices from E22 $\Delta$ A $\beta$  transgenic mice. This is opposed to a study which showed that synthetic E22 $\Delta$  A $\beta$  caused loss of synaptophysin signals in stained slice

cultures [22], although treatment with 1  $\mu$ M synthetic A $\beta$ 42 did not cause a significant reduction in the synaptophysin signal in that study. Further, primary cultures transfected with E693 $\Delta$  APP showed a loss of dendritic spine numbers compared to wt APP transfected cells [23]. These discrepancies to our data may be explained by the different techniques used. As discussed above, synthetic and recombinant A $\beta$  have different aggregation kinetics probably leading to different types of aggregates. Further, lipofected primary neurons strongly differ from transgenic slice cultures due to differential expression levels, time-course of expression and different properties of tissue and dissociated cultures (see below). Third, our transgenic models bear the APP<sub>swe</sub> mutation in addition to the intra-A $\beta$  mutation, which increases total A $\beta$  production. Nevertheless, we did not observe any effect on spine density in cultures from E22 $\Delta$ A $\beta$  transgenic mice. In addition to the number of spines, spine morphology has been shown to be affected in AD models [24]. Primary neurons transfected with E693 $\Delta$  APP displayed a specific loss of mushroom-shaped spines, but not of stubby or thin spines [23]. A loss of mushroom spines was also observed in APP<sub>SDL</sub> transgenic slice cultures, which produce A $\beta$  with the E693Q (Dutch) mutation [15, 25], as well as in hippocampal neurons treated with wt A $\beta$  oligomers [26]. However, both studies also showed that loss of mushroom spines was accompanied by an increase in the fraction of stubby spines, indicating a transition from mushroom to stubby shape rather than a specific loss of mushroom spines. Nevertheless, a direct comparison of the different A $\beta$  mutants and their effect on spine morphology in the same experimental system should be performed in the future.

Beside the loss of synapses, massive neurodegeneration is characteristic for AD. Previous studies from our lab showed that both transgenically produced A $\beta$  and treatment with recombinant wt A $\beta$  induced neurodegeneration in slice cultures, which was prominent in the presence of human tau expression [13, 15]. Here, we showed that wt and E22G A $\beta$  induced a tau-dependent cell death. As tau was expressed using neurotrophic Sindbis virus, we conclude that the affected cells were primarily neurons. In contrast, E22 $\Delta$  A $\beta$  did not show induction of toxicity. The absence of neurotoxicity is likely due to the lack of induced tau phosphorylation by E22 $\Delta$  A $\beta$ . While wt A $\beta$ 42 and E22G A $\beta$ 42 massively increases AT8 phosphorylation of tau, treatment with E22 $\Delta$  A $\beta$ 42 did not elevate phospho-tau signals above baseline. The AT8 epitope (Ser202/Thr205) is a major phosphorylation site in AD and used to classify the degree of AD pathology into different Braak stages [27]. Increased phosphorylation, among others at the AT8 epitope, has been shown to confer toxic properties to tau [13, 15]. Our data is in line with studies from other labs in which synthetic wt A $\beta$ 42 caused a strong MTT reduction in N2a and IMR-32 cells, while E22 $\Delta$  A $\beta$ 42 only caused a slight reduction in MTT signal [22]. Further, treatment of primary rat cortical cultures with synthetic wt and E22P A $\beta$ 42 but not E22 $\Delta$  A $\beta$ 40 or A $\beta$ 42 reduced cell survival [28]. Ovchinnikova and colleagues compared the toxicity of recombinant wt and E22 $\Delta$  A $\beta$ 40 and A $\beta$ 42 [9]. When added to primary neuronal cultures wt A $\beta$ 42 strongly induced LDH release, while E22 $\Delta$  A $\beta$ 42 did not affect cell survival. Interestingly, E22 $\Delta$  A $\beta$ 40 showed a detectable cell toxicity, although not as strong as wt A $\beta$ 42. The reason why we did not observe cell toxicity of E22 $\Delta$  A $\beta$ 40 may be explained by the use of a lower concentration (1  $\mu$ M compared to 7.4  $\mu$ M) and the use of an ex vivo tissue culture model. From our experience primary dissociated neuronal cultures are generally more susceptible to toxic insults than tissue cultures. In contrast to the lack of tau phosphorylation and tau-dependent cell death in our study, Umeda and colleagues found exacerbated tau pathology in APP<sub>osk</sub>-hTau double transgenic mice [29], suggesting that *in vivo* E22 $\Delta$  A $\beta$  may well affect tau biology. However, the study did not compare APP<sub>osk</sub>-hTau transgenic mice to APP<sub>wt</sub>-hTau transgenic mice or APP<sub>arc</sub>-hTau transgenic mice. Although speculative, APP<sub>wt</sub>-hTau or APP<sub>arc</sub>-hTau mice may have shown a stronger tau pathology than APP<sub>osk</sub>-hTau.

Taken together, we observed synaptic loss, tau phosphorylation and neuronal degeneration induced by wt and E22G A $\beta$ , both in a transgenic model and in a recombinant form, while E22 $\Delta$  A $\beta$  did not display any effects. Nevertheless, the lack of toxicity of E22 $\Delta$  A $\beta$  in our model requires further explanation. We showed differential aggregation properties of recombinant E22 $\Delta$  A $\beta$  in our previous study [8]. The type of E22 $\Delta$  A $\beta$  oligomers which cause neuronal degeneration and synaptic loss in AD patients with the Osaka APP mutations may either not be present in our model (see also discussion above on aggregation propensity of recombinant E22 $\Delta$  A $\beta$ ) or it may appear in the wrong localization. It has been speculated that E22 $\Delta$  A $\beta$  oligomers predominantly accumulate intra-neuronally and exert their toxic effects from inside the cell [30]. In line, Kondo and colleagues found intracellular A $\beta$  oligomers in induced pluripotent stem cell-derived neurons from E693 $\Delta$  AD patients but not in patients with the V717F APP mutation [31]. They hypothesized, that the pathophysiologic heterogeneity in AD may be due to the differential presence of intracellular A $\beta$  oligomers. Hence, familial AD caused by E693 $\Delta$  would belong to the intraneuronal-type of AD. E22 $\Delta$ A $\beta$  tg mice, used in our study, display intraneuronal A $\beta$  aggregates, associated with cognitive deficits, at 3 months of age [8]. However, as our slice cultures were prepared from P6-8 mice and cultured for 2 weeks, we do not expect the presence of intracellular A $\beta$  accumulation, which may explain the lack of toxicity in this model. Interestingly, arcA $\beta$  (E22G) transgenic mice also accumulate intraneuronal A $\beta$ , although at a later time point than E22 $\Delta$ A $\beta$  tg mice. Nevertheless, we previously showed that wt and E22G A $\beta$  toxicity depends on NMDA receptor signalling and that treatment of arcA $\beta$  transgenic cultures with anti-A $\beta$  antibodies rescued A $\beta$  toxicity, clearly indicating that both wt and E22G A $\beta$  at least have an extracellular phase in their toxicity mechanisms [13, 15]. Collectively, our findings therefore support the hypothesis of an intraneuronal toxicity mechanism by E22 $\Delta$ A $\beta$ .

### Pitfalls/limitations

We are aware that the main conclusion of our study, that E22G and E22 $\Delta$  A $\beta$  work via different mechanisms caused by differential aggregation properties and potential different localization of the aggregates, is only shown indirectly. A direct measure would be the analysis of A $\beta$  aggregates and their localization in the slice culture model. However, our study was affected by a limited access to animals and tissue, that did not allow for further analysis A $\beta$  aggregates or localization. Nevertheless, we base our conclusions on a combination of data obtained in the present study and of data from previous publications, where we showed differential properties and aggregation kinetics of wt, E22G and E22 $\Delta$  A $\beta$  in vitro and in vivo [8]. Thus, we are convinced that the conclusion drawn here are valid, however with taking the above-mentioned limitations into account.

### Supporting information

**S1 Table. P-values, mean differences and significances for all experiments.**  
(PDF)

**S1 Fig. Comparison of spine loss and cell toxicity between the 40- and 42-residue variants of recombinant wt, E22G and E22 $\Delta$  A $\beta$ .** These graphs summarize the findings of Figs 2B and 3B and were plotted to better compare the effects of wt A $\beta$ 42, E22G A $\beta$ 42 and E22 $\Delta$  A $\beta$ 42 to the respective 40-residue variant, wt A $\beta$ 40, E22G A $\beta$ 40 and E22 $\Delta$  A $\beta$ 40. A: Spine counts per  $\mu$ m dendrite. Spine density reduction after treatment with wt A $\beta$ 42 and E22G A $\beta$ 42 was higher compared to wt A $\beta$ 40 and E22G A $\beta$ 40, respectively. No difference was observed between E22 $\Delta$  A $\beta$ 42 and E22 $\Delta$  A $\beta$ 40. n = 10–15 B: Cytotoxicity in human tau-expressing slices

from wild-type mice treated with 1  $\mu$ M recombinant A $\beta$ . Wt A $\beta$ 42 and E22 $\Delta$  A $\beta$ 42 induced more cytotoxicity than wt A $\beta$ 40 and E22G A $\beta$ 40. No difference was observed between E22 $\Delta$  A $\beta$ 42 and E22 $\Delta$  s1A $\beta$ 40.  $n = 8-10$ . Data are means  $\pm$  SD. Statistical significance was assessed by one-way ANOVA with Tukey's multiple comparison test (\*\* $p < 0.01$ , \*\*\* $p < 0.001$ ). (TIF)

**S1 File.**  
(PDF)

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## Author Contributions

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